- ANSWER 4 OF 10 SCISEARCH COPYRIGHT (c) 2004 The Thomson Corporation. on 1.7 STN
- AN96:544888 SCISEARCH
- The Genuine Article (R) Number: UX582 GΑ
- AFRICAN-GREEN MONKEY KIDNEY (VERO) CELLS PROVIDE AN ALTERNATIVE HOST-CELL ΤI SYSTEM FOR INFLUENZA-A AND INFLUENZA-B VIRUSES
- GOVORKOVA E A; MURTI G; MEIGNIER B; DETAISNE C; WEBSTER R G (Reprint) ΑU
- ST JUDE CHILDRENS HOSP, DEPT VIROL & MOLEC BIOL, 332 N LAUDERDALE ST, CS MEMPHIS, TN, 38105 (Reprint); ST JUDE CHILDRENS HOSP, DEPT VIROL & MOLEC BIOL, MEMPHIS, TN, 38105; DI IVANOVSKII INST VIROL, MOSCOW 123098, RUSSIA; PASTEUR MERIEUX, F-69280 MARCY LETOILE, FRANCE; UNIV TENNESSEE, DEPT PATHOL, MEMPHIS, TN, 38163
- CYA USA; RUSSIA; FRANCE
- JOURNAL OF VIROLOGY, (AUG 1996) Vol. 70, No. 8, pp. 5519-5524. SO ISSN: 0022-538X.
- DTArticle; Journal
- FS LIFE
- LA ENGLISH
- REC
- Reference Count: 32 The preparation of live, attenuated human influenza virus AΒ vaccines and of large quantities of inactivated vaccines after the emergence or reemergence of a pandemic influenza virus will require an alternative host cell system, because embryonated chicken eggs will likely be insufficient and suboptimal. Preliminary studies indicated that an African green monkey kidney cell line (Vero) is a suitable system for the primary isolation and cultivation of influenza A viruses (E. A. Govorkova, N. V. Kaverin, L. V. Gubareva, B. Meignier, and R. G. Webster, J. Infect. Dis. 172:250-253, 1995). We now demonstrate for the first time that Vero cells are suitable for isolation and productive replication of influenza B viruses and determine the biological and genetic properties of both influenza A and B viruses in Vero cells; additionally, we characterize the receptors on Vero cells compared, with those on Madin-Darby canine kidney (MDCK) cells. Sequence analysis indicated that the hemagglutinin of Vero cell-derived influenza B viruses was identical to that of MDCK-grown counterparts but differed from that of egg-grown viruses at amino acid positions 196 to 198. Fluorescenceactivated cell sorting analysis showed that although Vero cells possess predominantly alpha 2,3 galactose-linked sialic acid, they are fully susceptible to infection with either human influenza A or B viruses. Moreover, all virus-specific polypeptides were synthesized in the same proportions in Vero cells as in MDCK cells. Electron microscopic and immunofluorescence studies confirmed that infected Vero cells undergo the same morphological changes as do other polarized epithelial cells. Taken together, these results indicate that Vero cell lines could serve as an

alternative host system for the cultivation of influenza A and B

viruses, providing adequate quantities of either virus to meet the vaccine

*Computer-Aided Design Dogs Kidney Neoplasms *L-Lactate Dehydrogenase: ME, metabolism Lung Neoplasms Rats Software Tumor Cells, Cultured Volatilization 0 (Anesthetics, Inhalation); EC 1.1.1.27 (L-Lactate Dehydrogenase) MEDLINE on STN ANSWER 15 OF 17 93055224 MEDLINE PubMed ID: 1331151 Enhanced detection of respiratory viruses using the shell vial technique and monoclonal antibodies. Lee S H; Boutilier J E; MacDonald M A; Forward K R Department of Microbiology, Victoria General Hospital, Halifax, Nova Scotia, Canada. Journal of virological methods, (1992 Sep) 39 (1-2) 39-46. Journal code: 8005839. ISSN: 0166-0934. Netherlands Journal; Article; (JOURNAL ARTICLE) English Priority Journals 199212 Entered STN: 19930122 Last Updated on STN: 19970203 Entered Medline: 19921203 The shell vial technique using A549 and MDCK cells, coupled with the use of Bartels respiratory viral monoclonal antibodies, was evaluated initially for the detection of 28 previously isolated respiratory viruses. All viruses were recovered and correctly identified. The shell vial-monoclonal antibody technique was then evaluated for virus isolation from 338 respiratory specimens and compared with the conventional tube method. Both methods gave rise to a total of 83 virus isolates. Of these isolates, 68 (20.1%) were isolated and identified by the shell vial-monoclonal method; 60 (17.8%) were culture-positive by the conventional tube method; forty-five (13.3%) were positive by both methods. The shell vial-monoclonal antibody method yielded 12 isolates of influenza A, two isolates of parainfluenza type 3 and one each of parainfluenza types 1 and 3, which were missed by the conventional tube method, indicating the superior sensitivity and specificity of the shell vial-monoclonal antibody method (Chi-square analysis, P = 0.001) for the detection of these viruses. Of the 50 RSV isolates, 29 were detected by both methods and there were 21 discrepancies between the two methods. The shell vial-monoclonal antibody method also improved the turn-around time for the respiratory virus groups. Check Tags: Human Adenoviruses, Human: IM, immunology Adenoviruses, Human: IP, isolation & purification Animals *Antibodies, Monoclonal Cell Line Dogs Evaluation Studies Influenza A Virus,

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(FILE 'HOME' ENTERED AT 14:23:11 ON 22 DEC 2004)

	FILE 'MEDLI	NE	E' ENTERED AT 14:23:19 ON 22 DEC 2004
L1	2115	S	MADIN-DARBY CANINE KIDNEY
L2	243	S	L1 AND INFLUENZA
L3	4	S	L1 AND RSV
L4	423547	S	1997>PY>1995
L5	8	S	L2 AND L4

FILE 'SCISEARCH' ENTERED AT 14:30:38 ON 22 DEC 2004

195 S L2 L6

10 S L4 AND L6 . L7

- AN 1998:754949 CAPLUS
- DN 130:152470
- ED Entered STN: 02 Dec 1998
- TI Cell lines of pulmonary and non-pulmonary origin as tools to study the effects of house dust mite proteinases on the regulation of epithelial permeability
- AU Winton, H. L.; Wan, H.; Cannell, M. B.; Gruenert, D. C.; Thompson, P. J.; Garrod, D. R.; Stewart, G. A.; Robinson, C.
- CS Department of Pharmacology & Clinical Pharmacology, St George's Hospital Medical School, London, SW17 ORE, UK
- SO Clinical and Experimental Allergy (1998), 28(10), 1273-1285 CODEN: CLEAEN; ISSN: 0954-7894
- PB Blackwell Science Ltd.
- DT Journal
- LA English
- CC 15-9 (Immunochemistry)
- Allergenic and non-allergenic proteinases from house dust mites (HDM) AΒ cause loss of adhesion between airway epithelial cells that may result in a loss of functional cohesion between the cells and thus assist in allergen presentation. Improved cellular assay systems are needed to ascertain the mechanisms involved. The authors surveyed a series of epithelial cell lines (Calu-3, 16HBE14o-, NCI-H292 and A549 from human airways, and MDCK from dog kidney) and established their utility for studies of the effects of HDM proteinases from D. pteronyssinus on epithelial permeability. In addition, the authors developed an improved method for measuring changes in epithelial permeability induced by HDM proteinases and other provocants. The permeability of epithelial monolayer cultures to mannitol was calculated from measurements of clearance using a technique that permits math. estimation and reduction of non-cellular diffusional constraints. Permeability was studied under control conditions and after perturbation of monolayers with HDM proteinases (separated into serine- and cysteine-proteinase classes) or chelation of extracellular Ca2+. Fluorescent antibody staining was used to investigate whether the cells expressed tight junctions (staining of ZO-1), desmosomes (staining of desmoplakin) and zonulae adherentes (staining of E-cadherin). The Calu-3 line was identified as an airway cell line that expressed functional tight junctions, desmosomes and zonulae adherentes. Calu-3 monolayers exhibited a low clearance and permeability to mannitol, similar to that seen in the extensively characterized MDCK cell line. Clearance and permeability were significantly increased by treatment with either HDM proteinase fraction or by calcium chelation. 16HBE14o- cells also had a low permeability to mannitol under control conditions and expressed a similar repertoire of functional proteins from major intercellular junctions. In contrast, NCI-H292 and A549 cell lines were functionally deficient in tight junctions, although they did express desmosomes and zonulae adherentes to a greater extent. Epithelial permeability was a more appropriate and sensitive index of epithelial perturbation than was tracer clearance. These results suggest that the Calu-3 and 16HBE14o- cell lines are useful tools in studying the mechanism of HDM proteinases on airway epithelial cell function. HDM proteinases of both cysteine and serine mechanistic classes were found to perturb epithelial adhesion and function.
- ST house dust mite proteinase airway epithelium permeability
- IT Animal cell line

(16HBE14o-; house dust mite proteinase effects on permeability of airway epithelium cell lines)

IT Animal cell line

(Calu-3; house dust mite proteinase effects on permeability of airway epithelium cell lines)

(FILE 'HOME' ENTERED AT 08:05:27 ON 22 DEC 2004)

L	1	FILE	'MEDLINE' ENTERED AT 08:05:35 ON 22 DEC 2004 3554 S MDCK
_	2		3035 S A549
L	_		125 S H292
L	4		17 S L1 AND L2
L	5		1 S L1 AND L3
L	6		1 S L1 AND L2 AND L3
		FILE	'BIOSIS' ENTERED AT 08:17:35 ON 22 DEC 2004
L	7		5 S L1 AND L3
L	8		45 S L1 AND L2
L	9	*	1 S L1 AND L2 AND L3
		FILE	'CAPLUS' ENTERED AT 08:20:14 ON 22 DEC 2004
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(FILE 'HOME' ENTERED AT 13:42:21 ON 22 DEC 2004)

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L1		2115 S MADIN-DARBY CANINE KIDNEY
L2		6 S L1 AND A549
L3		0 S L1 AND H292
L4		0 S CCL-1848
L5		56 S HUMAN LUNG EPITHELIAL CELL LINE
L6		1 S L5 AND L1
L7		0 S L5 AND MDCK
L8		1 S MDCK AND H292
	FILE	'BIOSIS' ENTERED AT 13:50:13 ON 22 DEC 2004
L9		0 S L1 AND L5
L10		23 S L1 AND A549
L11		3 S L1 AND H292

FILE 'MEDLINE' ENTERED AT 13:53:25 ON 22 DEC 2004

WEST Search History

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	L8	Madin-Darby canine kidney and RSV.clm.	4
	L7	Madin-Darby canine kidney.clm.	7
	L6	Madin-Darby canine kidney and influenza.clm.	48
	L5	Madin-Darby canine kidney and influenza	109
	L4	Madin-Darby canine kidney and RSV	33
	L3	Madin-Darby canine kidney	263
	L2	H292.clm.	7
	L1	H292	53

END OF SEARCH HISTORY







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			shell v J Clin N	rial techniqu Microbiol. 199	ie. 1 Mar;29(3)	cytial virus in 1 :463-5. ed for MEDLINE		ngeal se	ecretions by	/	

Related Articles, Links **8:** Johnston SL, Siegel CS. Evaluation of direct immunofluorescence, enzyme immunoassay, centrifugation culture, and conventional culture for the detection of respiratory syncytial virus. J Clin Microbiol. 1990 Nov;28(11):2394-7. PMID: 2254415 [PubMed - indexed for MEDLINE] Related Articles, Links 9: Olsen MA, Shuck KM, Sambol AR. Evaluation of Abbott TestPack RSV for the diagnosis of respiratory syncytial virus infections. Diagn Microbiol Infect Dis. 1993 Feb; 16(2):105-9. PMID: 8467621 [PubMed - indexed for MEDLINE] Related Articles, Links ☐ 10: Bartholoma NY, Forbes BA. Successful use of shell vial centrifugation and 16 to 18-hour immunofluorescent staining for the detection of influenza A and B in clinical specimens. Am J Clin Pathol. 1989 Oct;92(4):487-90. PMID: 2679041 [PubMed - indexed for MEDLINE] 11: Navarro-Mari JM, Sanbonmatsu-Gamez S, Perez-Ruiz M, De La Related Articles, Links Rosa-Fraile M. Rapid detection of respiratory viruses by shell vial assay using 'simultaneous culture of HEp-2, LLC-MK2, and MDCK cells in a single vial. J Clin Microbiol. 1999 Jul;37(7):2346-7. Erratum in: J Clin Microbiol 1999 Oct;37 (10):3436.PMID: 10364611 [PubMed - indexed for MEDLINE] Related Articles, Links ☐ 12: Ray CG, Minnich LL. Efficiency of immunofluorescence for rapid detection of common respiratory viruses. J Clin Microbiol. 1987 Feb;25(2):355-7. PMID: 3029168 [PubMed - indexed for MEDLINE] ☐ 13: Brinker JP, Doern GV. Related Articles, Links A comparison of commercially available monoclonal antibodies for direct and indirect immunofluorescence culture confirmation and direct detection of parainfluenza viruses. Diagn Microbiol Infect Dis. 1992 Nov-Dec; 15(8):669-72. PMID: 1335862 [PubMed - indexed for MEDLINE] ☐ 14: Hierholzer JC, Bingham PG, Coombs RA, Johansson KH, Related Articles, Links Anderson LJ, Halonen PE. Comparison of monoclonal antibody time-resolved fluoroimmunoassay with monoclonal antibody capture-biotinylated detector enzyme immunoassay for respiratory syncytial virus and parainfluenza virus antigen detection. J Clin Microbiol. 1989 Jun;27(6):1243-9. PMID: 2546973 [PubMed - indexed for MEDLINE] 15: Barenfanger J, Drake C, Mueller T, Troutt T, O'Brien J, Guttman Related Articles, Links R-Mix cells are faster, at least as sensitive and marginally more costly than

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	conventional cell lines for the detection of respi J Clin Virol. 2001 Aug;22(1):101-10. PMID: 11418357 [PubMed - indexed for MEDLINE]	ratory vii	ruses.		
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	Evaluation of a monoclonal antibody pool for rarespiratory viral infections. J Clin Microbiol. 1989 Mar;27(3):448-52. PMID: 2541165 [PubMed - indexed for MEDLINE]	apid diag	nosis of		
□17:	Klespies SL, Cebula DE, Kelley CL, Galehouse D, Maur	er CC.	Related A	rticles,	Links
	Detection of enteroviruses from clinical specimshell vial culture and monoclonal antibody assar J Clin Microbiol. 1996 Jun;34(6):1465-7. PMID: 8735099 [PubMed - indexed for MEDLINE]		oin ampl	ificatio	on
□18:	Shih SR, Tsao KC, Ning HC, Huang YC, Lin TY.		Related A	Articles,	Links
	Diagnosis of respiratory tract viruses in 24 h by staining of shell vial cultures containing Madin-(MDCK) cells. J Virol Methods. 1999 Aug;81(1-2):77-81. PMID: 10488764 [PubMed - indexed for MEDLINE]				
□ 19:	Alexander R, Lamb D, White D, Wentzel T, Politis S, Ri J, van Ruyven D, Kelly N, Garland SM.	<u>insburger</u>	Related A	Articles,	Links
	'RETCIF': a rapid, sensitive method for detection large numbers of clinical samples. J Virol Methods. 2001 Sep;97(1-2):77-85. PMID: 11483219 [PubMed - indexed for MEDLINE]	on of viru	ses, app	licable	e for
□ 20:	Van Doornum GJ, De Jong JC.		Related A	Articles,	Links
	Rapid shell vial culture technique for detection adenoviruses in fecal specimens: comparison w isolation method. J Clin Microbiol. 1998 Oct;36(10):2865-8. PMID: 9738034 [PubMed - indexed for MEDLINE]				
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Dec 13 2004 14:18:14

WEST Search History

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	L1	MDCK and A549 and H292	12

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	L7	Madin-Darby canine kidney.clm.	7
	L6	Madin-Darby canine kidney and influenza.clm	. 48
	L5	Madin-Darby canine kidney and influenza	109
	L4	Madin-Darby canine kidney and RSV	33
⊡	L3	Madin-Darby canine kidney	263
	L2	H292.clm.	7
	L1	H292	53

END OF SEARCH HISTORY

Hit List

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Search Results - Record(s) 1 through 4 of 4 returned.

☐ 1. Document ID: US 6573080 B2

L2: Entry 1 of 4

File: USPT

OH

Jun 3, 2003

US-PAT-NO: 6573080

DOCUMENT-IDENTIFIER: US 6573080 B2

TITLE: Mixed cell diagnostic systems

DATE-ISSUED: June 3, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Scholl; David R. Athens
Huang; Yung T. Richmond Heights

Goodrum; Patricia Gail Ray Athens Ol

US-CL-CURRENT: 435/235.1; 435/325, 435/5, 435/6, 435/8

Full Title Citation Front Review Classification Date Reference

☐ 2 Document ID: US 6406842 B2

L2: Entry 2 of 4 File: USPT Jun 18, 2002

US-PAT-NO: 6406842

DOCUMENT-IDENTIFIER: US 6406842 B2

TITLE: Mixed cell diagnostic systems

DATE-ISSUED: June 18, 2002

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Scholl; David R. Athens OH
Huang; Yung T. Richmond Heights OH
Goodrum; Patricia Gail Ray Athens OH

US-CL-CURRENT: 435/5; 435/235.1, 435/325, 435/8

Full Title Citation Front Review Classification Date Reference (1997) (1997) (1997) (1997) (1997)

☐ 3. Document ID: US 6376172 B1

L2: Entry 3 of 4

File: USPT

Apr 23, 2002

US-PAT-NO: 6376172

DOCUMENT-IDENTIFIER: US 6376172 B1

TITLE: Mixed cell diagnostic systems

DATE-ISSUED: April 23, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE COUNTRY

Scholl; David R.

Athens

OH

Huang; Yung T.

Richmond Heights

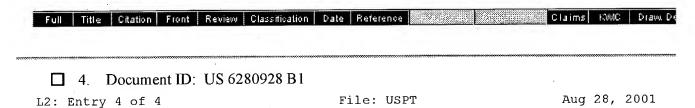
OH

Goodrum; Patricia Gail Ray

Athens

OH

US-CL-CURRENT: 435/5; 435/235.1, 435/325, 435/8



US-PAT-NO: 6280928

DOCUMENT-IDENTIFIER: US 6280928 B1

TITLE: Mixed cell diagnostic systems

DATE-ISSUED: August 28, 2001

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE COUNTRY

Scholl; David R.

Athens

OH

Huang; Yung T.

Richmond Heights

Goodrum; Patricia Gail Ray

Athens

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US-CL-CURRENT: 435/5; 435/235.1, 435/325, 435/8

